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Annex US.II, page 1

NATIONAL CHAPTER - US

FORM PTO-1390 (REV 11-2000)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER SHIM1130
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				U.S. APPLICATION NO. (If known, see 37 CFR 1.5)
INTERNATIONAL APPLICATION NO. PCT/JP00/06987		INTERNATIONAL FILING DATE 6 October 2000		10/089789
PRIORITY DATE CLAIMED 6 October 1999				
TITLE OF INVENTION CARBONYL STRESS-DECREASING AGENT				
APPLICANT(S) FOR DO/EO/US Toshio MIYATA and Kiyoshi KUROKAWA				
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:				
<p>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 USC 371.</p> <p>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing 35 USC 371.</p> <p>3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 USC 371(f)). The submission must include items (5), (6), (9), and (21) indicated below.</p> <p>4. <input type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31).</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)) <ul style="list-style-type: none"> a. <input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau). b. <input checked="" type="checkbox"/> has been communicated by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US) </p> <p>6. <input checked="" type="checkbox"/> An English language translation of the International Application as filed (35 USC 371(c)(2)) <ul style="list-style-type: none"> a. <input checked="" type="checkbox"/> is attached hereto b. <input type="checkbox"/> has been previously submitted under 35 USC 154(d)(4) </p> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ul style="list-style-type: none"> a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> have been communicated by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made. </p> <p>8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 USC 371(c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An unsigned oath or declaration of the inventor(s) (35 USC 371(c)(4)).</p> <p>10. <input type="checkbox"/> An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 USC 371(c)(5)).</p>				
Items 11. to 16. below concern other document(s) or information included:				
<p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input checked="" type="checkbox"/> A FIRST preliminary amendment.</p> <p>14. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>15. <input type="checkbox"/> A substitute specification.</p> <p>16. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>17. <input type="checkbox"/> A computer-readable form of sequence listing in accordance with PCT Rule 13 and 35 USC 1821-1825</p> <p>18. <input type="checkbox"/> A second copy of the published International Application under 35 USC 154(d)(4).</p> <p>19. <input type="checkbox"/> A second copy of the English language translation of the International Application under 35 USC 154(d)(4).</p> <p>20. <input checked="" type="checkbox"/> Other items or information: Check No. 505102 in the amount of \$1300.00; Return Postcard</p>				
Express Mail Label No.: EV047297082				

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NATIONAL CHAPTER - US

Annex US.II, page 2

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)		INTERNATIONAL APPLICATION NO. PCT/JP99/05789		ATTORNEY'S DOCKET NO. SHIM1120	
21. <input checked="" type="checkbox"/> The following fees are submitted:				CALCULATIONS	PTO USE ONLY
BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5)): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2) paid to USPTO and International Search Report not prepared by the EPO or JPO International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by EPO or JPO International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4). International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4).				\$1,040.00	
				\$ 890.00	
				\$ 740.00	
				\$ 710.00	
				\$ 100.00	
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$ 890.00	
Surcharge of \$130.00 for furnishing the oath or declaration later than <u>X</u> 20 ___ 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				\$130.00	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total Claims	18 - 20 =	0	x \$ 18.00	\$	
Independent Claims	2 - 3 =	0	x \$ 84.00	\$	
Multiple dependent claim(s) (if applicable)				+\$280.00	\$ 280.00
TOTAL OF ABOVE CALCULATIONS =				\$ 1020.00	
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.				\$	
SUBTOTAL =				\$ 1300.00	
Processing fee of \$130. for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)). +				\$	
TOTAL NATIONAL FEE =				\$1300.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	
TOTAL FEES ENCLOSED =				\$ 1300.00	
				Amount to be refunded:	\$
				charged:	\$
a. <input checked="" type="checkbox"/> A check in the amount of \$ <u>1300.00</u> to cover the above fees is enclosed. b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>50-1355</u> . A duplicate copy of this sheet is enclosed.					
NOTE: Where an appropriate timelimit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO: GRAY CARY WARE & FREIDENRICH 4365 Executive Drive, Suite 1100 San Diego, California 92121-2133					
 LISA A. HAILE, Reg. No. 38,347 April 2, 2002 Date					
PHONE: 858/677-1456 - FAX: 858/677-1465					

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PATENT
Attorney Docket No.: SHIM1130

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Miyata and Kurokawa
Serial No.: Not yet assigned
Filed: April 2, 2002
Based on: PCT/JP00/06987
Internat'l. Priority
Filing Date: October 6, 2000 Date: October 6, 1999
Title: CARBONYL STRESS-DECREASING AGENT

BOX PCT
Commissioner for Patents
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

In connection with the above-identified patent application, entry of the amendments and consideration of the following remarks are respectfully requested.

AMENDMENTS

In the Claims

Please add the following new claims:

4. The carrier of claim 3, wherein the carrier is selected from the group consisting of: synthetic or naturally-occurring organic macro-molecular compounds; inorganic materials, such as glass beads, silica gel, alumina, and activated charcoal; and materials coated with polysaccharide(s) or synthetic polymer(s) thereof.

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I HEREBY CERTIFY THAT THIS PAPER OR FEE IS BEING DEPOSITED WITH THE UNITED STATES POSTAL SERVICE "EXPRESS MAIL POST OFFICE TO ADDRESSEE" SERVICE UNDER 37 C.F.R. 1.10 ON THE DATE INDICATED ABOVE AND IS ADDRESSED TO:	
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In re Application of
Miyata and Kurokawa
Application No.: Not yet assigned
Filed: April 2, 2002
Based on International Appl. No. PCT/JP00/06987
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5. The carrier of claim 3, wherein the shape of the carrier is selected from the group consisting of: membranous, fibrous, granular-shaped, hollow fiber-like, non-woven fabric-like, porous, and honeycomb-shaped carrier.

6. The carrier of claim 3, wherein the carrier has a body fluid contact area that can be controlled by changing: thickness, surface area, diameter, length, shape, and/or size of the carrier.

7. The carrier of claim 3, wherein the carrier is a hemodialysis membrane.

8. The carrier of claim 3, wherein the biguanide agent is immobilized on the carrier by physical adsorption, a specific biochemical binding reaction, ion binding, covalent bonding, or grafting.

9. An adsorbent of carbonyl compounds comprising any of the carriers of claim 3 to 8.

10. A peritoneal dialysate solution comprising the carbonyl stress-decreasing agent of claim 1.

11. A method for removing carbonyl compounds comprising the step of contacting the carrier of claim 3 with a body fluid selected from the group consisting of: blood, blood plasma and peritoneal dialysate.

12. The method of claim 11, wherein the removal of the carbonyl compounds are carried out during in vivo or ex vivo blood purification step.

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13. The method of claim 12, wherein the blood purification step comprises one or more steps selected from the group consisting of: hemodialysis, blood filtration, blood filtration dialysis, blood adsorption, and blood plasma separation.

In the Description

Page 4 line 1 to line 12, please change as follows:

- [1] a carbonyl stress-decreasing agent comprising a biguanide agent or pharmacologically acceptable salt thereof as an active ingredient;
- [2] the carbonyl stress-decreasing agent of [1], wherein the biguanide agent is a compound selected from the group consisting of: phenformin, metformin, buformin, and pharmacologically acceptable salts thereof;
- [3] a carrier, on which a biguanide agent has been immobilized;
- [4] the carrier of [3], wherein the carrier is selected from the group consisting of: synthetic or naturally-occurring organic macro-molecular compounds; inorganic materials, such as glass beads, silica gel, alumina, and activated charcoal; and materials coated with polysaccharide(s) or synthetic polymer(s) thereof;
- [5] the carrier of [3], wherein the shape of the carrier is selected from the group consisting of: membranous, fibrous, granular-shaped, hollow fiber-like, non-woven fabric-like, porous, and honeycomb-shaped carrier;
- [6] the carrier of [3], wherein the carrier has a body fluid contact area that can be controlled by changing: thickness, surface area, diameter, length, shape, and/or size of the carrier;
- [7] the carrier of [3], wherein the carrier is a hemodialysis membrane;
- [8] the carrier of [3], wherein the biguanide agent is immobilized on the carrier by physical adsorption, a specific biochemical binding reaction, ion binding, covalent bonding, or grafting;

In re Application of
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- [9] an adsorbent of carbonyl compounds comprising any of the carriers of [3] to [8];
[10] a peritoneal dialysate solution comprising the carbonyl stress-decreasing agent of [1];
[11] a method for removing carbonyl compounds comprising the step of contacting the carrier of [3] with a body fluid selected from the group consisting of: blood, blood plasma and peritoneal dialysate;
[12] the method of [11], wherein the removal of the carbonyl compounds are carried out during in vivo or ex vivo blood purification step; and
[13] the method of [12], wherein the blood purification step comprises one or more steps selected from the group consisting of: hemodialysis, blood filtration, blood filtration dialysis, blood adsorption, and blood plasma separation.

REMARKS

Upon entry of the present amendment, claims 1 to 13 will be pending.

In summary, the amendments and new claims address formalities and are supported by the claims as originally filed, and, therefore, do not add new matter. Accordingly, entry of the amendments and new claims is respectfully requested.

It is submitted that the claims are in condition for allowance, and a notice to that effect respectfully is requested. The Examiner is invited to contact Applicants' undersigned representative if there are any questions relating to this application.

In re Application of
Miyata and Kurokawa
Application No.: Not yet assigned
Filed: April 2, 2002
Based on International Appl. No. PCT/JP00/06987
International Filing Date: 6 October 2000
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PATENT
Attorney Docket No.: SHIM1130

Please charge any additional fees, or make any credits, to Deposit Account No. 50-1355.

Respectfully submitted,

Date: April 2, 2002



Lisa A. Haile, Ph.D.
Reg. No. 38,347
Attorney for Applicant
Telephone No.: (858) 677-1456
Facsimile No.: (858) 677-1465

GRAY CARY WARE & FREIDENRICH LLP
4365 Executive Drive, Suite 1100
San Diego, California 92121-2133

US Customer No. 28213

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DESCRIPTION

CARBONYL STRESS-DECREASING AGENT

5 Technical Field

The present invention relates to carbonyl stress-decreasing agents.

Background Art

10 An enhanced state of *in vivo* carbonyl compound production by non-enzymatic biochemical reactions is called "carbonyl stress". Carbonyl compounds are considered to be involved in aging and adult diseases, such as diabetes mellitus, and arteriosclerosis, via the Maillard reaction. The Maillard reaction is a non-enzymatic
15 glycation reaction between a reducing sugar, such as glucose, and amino acids or proteins. Maillard reported this reaction in 1912, focusing on a phenomenon of brown coloration arising upon heating a mixture consisting of amino acid and reducing sugar (Maillard, L. C., Compt. Rend. Soc. Biol., 72: 599 (1912)). The Maillard reaction
20 is involved in brown coloration, generation of aromatic components, taste and protein denaturation, and such, during heating or storage of foods. Therefore, this reaction has been mainly studied in the field of food chemistry.

However, in 1968, glycated hemoglobin (HbA_{1c}), a micro fraction
25 of hemoglobin, was identified *in vivo*, and was further demonstrated to increase in patients with diabetes (Rahbar. S., Clin. Chim. Acta, 22: 296 (1968)). These findings brought attention to the significance of *in vivo* Maillard reactions, and the relationship between the reaction, the onset of adult diseases, such as diabetic
30 complications and arteriosclerosis, and the progress of aging. For example, pyrraline and pentosidine, the late-stage products formed at post-Amadori compound formation reaction stages (advanced glycation end products; hereinafter abbreviated as AGE), are considered to serve as indices of aging and diabetes mellitus. In
35 fact, highly reactive carbonyl compounds and AGE are accumulated at very high levels in blood and tissues of chronic renal failure patients,

regardless of the presence or absence of hyperglycemia (Miyata, T. et al., Kidney Int., 51:1170-1181, 1997; Miyata, T. et al., J. Am. Soc. Nephrol., 7:1198-1206, 1996; Miyata, T. et al., Kidney Int. 55:389-399, 1999; Miyata, T. et al., J. Am. Soc. Nephrol. 9:2349-2356, 1998). This accumulation is ascribed to carbonyl stress in renal failure, which modifies proteins as a result of Maillard reaction with carbonyl compounds derived from sugars and lipids with amino groups (Miyata, T. et al., Kidney Int. 55:389-399, (1999)).

Thus, improving the carbonyl stress state by removing carbonyl compounds, which are generated *in vivo*, can result in the suppression of AGE formation associated with renal failure and thereby reduce tissue damages.

In peritoneal dialysis, waste products are excreted from blood across the peritoneum to the peritoneal dialysate. A peritoneal dialysate with high osmotic pressure (containing glucose, icodextrin, amino acids, and so on) delivers highly reactive carbonyl compounds accumulated in blood of renal failure patients across the peritoneum into the peritoneal dialysate in peritoneal cavity. This results in an increase in carbonyl compound concentration within the peritoneal dialysate to cause a carbonyl stress state. As a result, the peritoneal function is lowered, due to the modification of intraperitoneal proteins with carbonyl; this reaction, in turn, is presumed to be involved in the impairment of water-removing ability and ingraevescence of peritoneal sclerosis (Miyata, T. et al., Kidney Int., 58:425-435, 2000; Inagi R., et al., FEBS Lett., 463:260-264, 1999; Ueda, Y., et al., Kidney Int. (in press); Combet, S., et al., J. Am. Soc. Nephrol., 11:717-728, 2000).

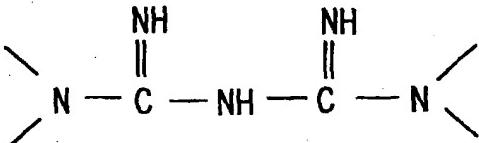
Indeed, the intraperitoneal carbonyl stress state induced by the introduction of glucose in peritoneal dialysis patients was demonstrated by immunohistochemical examination of the endothelia and mesothelia (Yamada, K. et al., Clin. Nephrol., 42: 354-361, 1994; Nakayama, M. et al., Kidney Int., 51: 182-186, 1997; Miyata, T. et al., Kidney Int., 58:425-435, 2000; Inagi R., et al., FEBS Lett., 463:260-264, 1999; Combet, S., et al., J. Am. Soc. Nephrol., 11:717-728, 2000). Thus, the carbonyl stress is also presumed to cause morphorgical changes in the peritoneum accompanied by

functional (water-removing ability) impairment in dialysis patients. Therefore, a method to decrease the stress is needed in the art.

As a method for decreasing carbonyl stress in peritoneal dialysis patients, the present inventor has filed a patent (PCT/JP99/04521) relating to the use of carbonyl compound-trapping agents, such as 5 aminoguanidine.

Biguanide agents, which have been used as therapeutic agents for diabetes mellitus, antimicrobial drugs, and antimalarial drugs, are known as compounds with a guanidine backbone. The biguanide 10 agents have a basal backbone shown in formula (1), and contain a highly reactive imino group (=NH). Thus, the biguanide agents are expected to have a carbonyl compound-removing activity, like those of aminoguanidine. However, it has been believed that biguanide agents lack the glycation-suppressing effect because of the difference in 15 three-dimensional structure between the two compounds ("New chemical therapy for diabetes mellitus" pp 22-31, 3. Biguanide Agents; S Tanaka, Medical Core (1997)).

(1)



20

Disclosure of the Invention

The object of the present invention is to provide carbonyl stress-decreasing agents that are effective against the state of systemic carbonyl stress.

25 The present inventor searched for compounds that remove carbonyl compounds accumulated in blood and such. As described in the reference of prior art cited above, the biguanide agents have been believed to lack the glycation-suppressing activity. However, the present inventor found that the concentration of carbonyl 30 compounds in a medium can be reduced by using a biguanide agent, such as phenformin, metformin, and buformin, and thus, completed the present invention. More specifically, the present invention relates to the following :

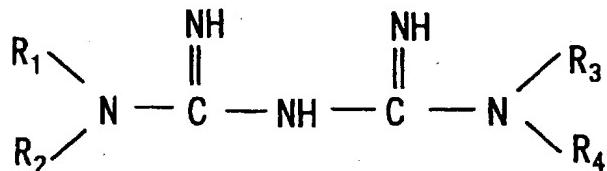
- [1] a carbonyl stress-decreasing agent comprising a biguanide agent or pharmacologically acceptable salt thereof as an active ingredient;
- [2] the carbonyl stress-decreasing agent of [1], wherein the biguanide agent is a compound selected from the group consisting of: phenformin,
5 metformin, and buformin; or pharmacologically acceptable salts thereof;
- [3] a carrier, on which a biguanide agent has been immobilized;
- [4] an adsorbent of carbonyl compounds, which comprises the carrier of [3]; and
- 10 [5] a method for removing carbonyl compounds, which comprises the step of contacting the carrier of [3] with patient's blood or peritoneal dialysate.

Further, the present invention relates to the use of biguanide agents or pharmacologically acceptable salts thereof to produce the carbonyl stress-decreasing agents. The present invention also relates to the use of carriers, on which a biguanide agent has been immobilized, to produce carbonyl-compound adsorbents.

Biguanide agents are a group of compounds which have a guanidine backbone; they have been used as therapeutic agents for diabetes mellitus. Recently, biguanide agents have been widely used in place of sulfonyl urea (SU) agent, an agent that had been used as an oral antidiabetic drug for a long period. It has been reported that biguanides lack the activity to enhance insulin secretion, but have the activity of enhancing glycolysis to reduce blood glucose levels
20 in living bodies. However, the interaction between biguanides and carbonyl compounds has not yet been reported, and thus, the finding of the present invention is entirely new.

As used herein, the term "biguanide agent" refers to, for example, a compound having the structure of formula (1) shown below.
30 Wherein, R₁, R₂, R₃, and R₄ are independently selected from H, phenyl group, and alkyl group. The alkyl group includes phenetyl group, methyl group, butyl group, etc.

Formula (1) :



Compounds with the above structure that act on carbonyl compounds causing carbonyl stress to inhibit the protein-modifying action thereof are used in the present invention.

Causative carbonyl compounds of carbonyl stress of the present invention include, for example, the following compounds that accumulate in blood of renal failure patients due to oxidative stress:

Carbonyl compounds derived from carbohydrates:

- 10 • arabinose
- glyoxal
- methylglyoxal
- 3-deoxyglucosone.

Carbonyl compound derived from ascorbic acid:

- 15 • dehydroascorbic acid.

Carbonyl compound derived from lipid:

- hydroxynonenal
- malondialdehyde
- acrolein.

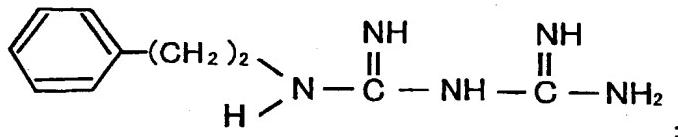
20 Further, during sterilization or storage of peritoneal dialysate, for example, the following carbonyl compounds are known to be produced in the peritoneal dialysate (Richard, J. U. et al., Fund. Appl. Toxic., 4: 843-853 (1984)). These carbonyl compounds, which are transferred through dialysis into the body of patients, 25 are also pointed out as the cause of the carbonyl stress state in patients:

- 3-deoxyglucosone
- 5-hydroxymethylfurfural
- formaldehyde
- 30 • acetaldehyde
- glyoxal
- methylglyoxal
- levulinic acid

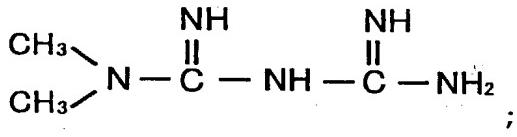
- furfural
- arabinose.

A carbonyl stress-decreasing agent of the present invention comprises a biguanide agent as the active ingredient. As used herein, the term "decreasing carbonyl stress" refers to reducing the reactivity of carbonyl compounds in a medium which is contacted with living bodies, and thus, alleviating protein modification. Specifically, for example, when a compound has a potential for adsorbing or removing carbonyl compounds, or a potential for reducing the reactivity to amino groups, then the compound is assumed to have the activity of relieving carbonyl stress. The term "medium which is contacted with living bodies" specifically refers to peritoneal dialysate, blood, and other body fluids. Biguanide agents that can be used in the present invention include the following compounds, and pharmacologically acceptable salts thereof:

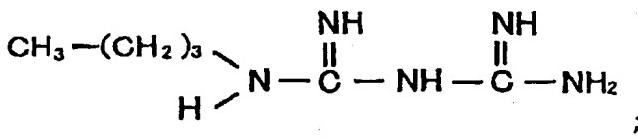
phenformin (phenethyl biguanide)



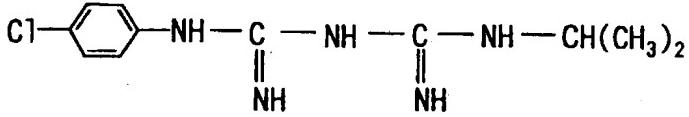
metformin (dimethyl biguanide)



buformin (butyl biguanide)

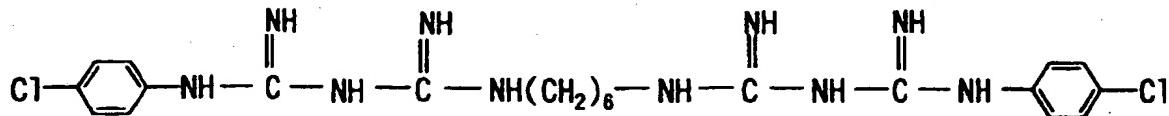


proguanil



; and

chlorhexidine



A carbonyl stress-decreasing agent of the present invention can be used in combination with compounds comprising, for example, the 5 following compounds, or derivatives thereof, that function as a carbonyl compound-trapping agents. The term "derivatives" herein, refers to compounds having an atomic or molecular substitution(s) at any position of the parent compound.

- (1) Guanidine derivatives, such as methylguanidine (JP-A Sho 10 62-142114; JP-A Sho 62-249908; JP-A Hei 1-56614; JP-A Hei 1-83059; JP-A Hei 2-156; JP-A Hei 2-765; JP-A Hei 2-42053; JP-A Hei 6-9380; and Published Japanese Translation of International Publication 5-505189);
- (2) hydrazine derivatives, such as sulfonylhydrazine;
- (3) five-membered heterocyclic compounds having two nitrogen atoms, such as pyrazolone (JP-A Hei 6-287179), pyrazoline (JP-A Hei 10-167965), pyrazole (JP-A Hei 6-192089; JP-A Hei 6-298737; and JP-A Hei 6-298738), imidazolidine (JP-A Hei 5-201993; JP-A Hei 6-135968; JP-A Hei 7-133264; and JP-A Hei 10-182460), and hydantoin (JP-A Hei 20 6-135968);
- (4) five-membered heterocyclic compounds having three nitrogen atoms, such as triazole (JP-A Hei 6-192089);
- (5) five-membered heterocyclic compounds having a nitrogen atom and a sulfur atom, such as thiazoline (JP-A Hei 10-167965), thiazole (JP-A Hei 4-9375; and JP-A Hei 9-59258), and thiazolidine (JP-A Hei 25 5-201993; JP-A Hei 3-261772; JP-A Hei 7-133264; and JP-A Hei 8-157473);
- (6) five-membered heterocyclic compounds having a nitrogen atom and an oxygen atom, such as oxazole (JP-A Hei 9-59258);
- (7) nitrogen-containing six-membered heterocyclic compounds, such as pyridine (JP-A Hei 10-158244; JP-A Hei 10-175954), and pyrimidine (Published Japanese Translation of International Publication 30 7-500811);

- (8) nitrogen-containing condensed heterocyclic compounds, such as indazole (JP-A Hei 6-287180), and benzimidazole (JP-A Hei 6-305964), quinoline (JP-A Hei 3-161441);
- (9) sulfur- and nitrogen-containing condensed heterocyclic compounds,
5 such as benzothiazole (JP-A Hei 6-305964);
- (10) sulfur-containing condensed heterocyclic compounds, such as benzothiophene (JP-A Hei 7-196498);
- (11) oxygen-containing condensed heterocyclic compounds, such as benzopyran (JP-A Hei 3-204874; and JP-A Hei 4-308586);
- 10 (12) nitrogenous compounds, such as carbazoyl (JP-A Hei 2-156; and JP-A Hei 2-753), carbazic acid (JP-A Hei 2-167264), and hydrazine (JP-A Hei 3-148220);
- (13) quinines, such as benzoquinone (JP-A Hei 9-315960), and hydroquinone (JP-A Hei 5-9114);
- 15 (14) aliphatic dicarboxylic acids (JP-A Hei 1-56614; and JP-A Hei 5-310565);
- (15) silicone containing compounds (JP-A Sho 62-249709);
- (16) organic germanium compounds (JP-A Hei 2-62885; JP-A Hei 5-255130; JP-A Hei 7-247296; and JP-A Hei 8-59485);
- 20 (17) flavonoids (JP-A Hei 3-240725; JP-A Hei 7-206838; JP-A Hei 9-241165; and WO 94/04520);
- (18) alkylamines (JP-A Hei 6-206818; JP-A Hei 9-59233; JP-A Hei 9-40626; and JP-A Hei 9-124471);
- (19) amino acids (Published Japanese Translation of International
25 Publication 4-502611; and Published Japanese Translation of International Publication 7-503713);
- (20) aromatic compounds, such as ascochlorin (JP-A Hei 6-305959), benzoic acid (WO 91/11997), and pyrrolo-naphthyridinium (JP-A Hei 10-158265);
- 30 (21) polypeptides (Published Japanese Translation of International Publication 7-500580);
- (22) vitamins, such as pyridoxamine (WO 97/09981);
- (23) SH group-containing compounds, such as glutathione, cysteine, and N-acetylcysteine;
- 35 (24) SH group-containing proteins, such as reduced albumin;
- (25) tetracyclines (JP-A Hei 6-256280);

- (26) chitosans (JP-A Hei 9-221427) ;
- (27) tannins (JP-A Hei 9-40519) ;
- (28) quaternary ammonium ion-containing compounds;
- (29) ion exchange resins; and
- 5 (30) inorganic compounds, such as activated carbon, silica gel, alumina, and calcium carbonate.

A carbonyl stress-decreasing agent of the present invention can be formulated in combination with physiologically acceptable carriers, excipients, diluents, and so on, to be administered orally or 10 parenterally. Dosage forms for oral drug include glandules, powders, tablets, capsules, solutions, emulsions, suspensions, etc.; and, those for parenteral drug include injections, drops, agents for external use, suppositories, and so on. The term "injection" encompasses intravenous injection, subcutaneous injection, 15 intramuscular injection, intraperitoneal injection, and such. The agents for external use include intranasal drugs, adhesive skin patches, ointments, and so on. Such dosage forms with a biguanide agent as the base can be formulated by conventional methods.

For example, tablets for oral administration can be produced 20 by combining biguanide agents with excipients, disintegrating agents, binding agents, lubricants, and such, to press-form the mixture. Typical excipients are exemplified by lactose, starch, mannitol, and the like. Calcium carbonate, calcium carboxymethyl cellulose, and such are generally used as disintegrating agents. Gum Arabic, 25 carboxymethyl cellulose, polyvinylpyrrolidone, and such are typically used as binding agents. Known lubricants include talc, magnesium stearate, and so on.

A tablet containing a carbonyl stress-decreasing agent of the present invention can be coated for masking or for making into 30 enteric-coated preparations according to conventional methods. Such coating agents include ethyl cellulose, polyoxyethyleneglycol, and so on.

An injection can be prepared by dissolving a biguanide agent 35 as the base together with an appropriate dispersant, or dissolving or dispersing the agent in a dispersion medium. Depending on the type of the selected dispersion medium, the dosage form can be an aqueous

solution or an oleaginous solution. Dispersion media used to prepare aqueous solutions includes distilled water, physiological saline, Ringers solution, and so on. Various vegetable oils, propyleneglycol, and the like can be exemplified as the dispersion media for preparing 5 an oleaginous solution. Additionally, preservatives, such as paraben, can also be added according to needs. Further, known isotonizing agents, such as sodium chloride, and glucose; and pH-adjusting agents, such as hydrochloric acid, and sodium hydroxide, can be added to the injection. Furthermore, soothing agents, such 10 as benzalkonium chloride, and procaine hydrochloride, can be added to the injection.

A carbonyl stress-decreasing agent of the present invention can be formulated for external use as a solid, liquid, or semi-solid composition comprising a biguanide agent. The solid or liquid 15 composition for external use can be prepared according to the method mentioned above for preparing compositions. The semi-solid composition can be prepared by adding a thickening agent(s) to an appropriate solvent(s) according to needs. Such solvents include water, ethyl alcohol, polyethyleneglycol, and so on. Typical 20 thickening agents include bentonite, polyvinyl alcohol, acrylic acid, methacrylic acid, polyvinylpyrrolidone, and so on. The composition may contain preservatives, such as benzalkonium chloride. Further, an agent of the present invention can also be formulated into a 25 suppository by the combined use of an oily base, such as cacao butter, or an aqueous gel base, such as cellulose derivatives, as a carrier.

The biguanide agents, used as the base of the carbonyl stress-relieving agents of the present invention, are compounds that have been previously used as pharmaceuticals. Accordingly, required 30 amounts of the agents, within a dose range ensuring the safety in usual cases, can be administered to mammals including human. The doses are properly selected depending on the type of administration method (dosage form), and condition of subject (size, age, sex, symptom, etc.). Generally, in the case of oral administration, a dose of typically 0.001 to 10mg, more preferably 0.01 to 1mg/day/kg body 35 weight (human adult) is administered to gain a carbonyl stress-decreasing effect. The frequency of administration can be

selected properly, for example, from a range of 1 to 5 times a day. No explicit toxicity is recognized for the biguanide agents of the present invention administered to decrease carbonyl stress.

A carbonyl stress-decreasing agent of the present invention can 5 be administered to a living body by adding the agent to a peritoneal dialysate. Peritoneal dialysis is carried out by injecting a dialysate into peritoneal cavity; thus, a carbonyl stress-decreasing agent of the present invention can be administered by previously adding the agent into the peritoneal dialysate. The carbonyl 10 compounds that exude into the peritoneal dialysate react with the biguanide agent to be detoxified, and, as a result, the carbonyl stress state is improved. Further, a biguanide agent previously added to the dialysate is expected to preventively detoxify the carbonyl 15 compounds produced by various treatments of the dialysate, such as autoclaving during the process of production, and storage. A biguanide agent is added at a concentration of, for example, 0.5 to 100 mM, typically 1 to 50 mM, and preferably 5 to 20 mM to a peritoneal dialysate of the present invention.

A carbonyl stress-decreasing agent of the present invention can 20 not only be directly administered to a living body, but also can be contacted with blood or dialysate outside the living body in methods for decreasing carbonyl stress. A biguanide agent is advantageously immobilized on a carrier for such methods.

There is no particular restriction on the carrier that is used 25 for immobilizing a biguanide agent of the invention, so long as the carrier is harmless to human body, and is sufficiently safe and stable as a material to be directly contacted with blood or peritoneal dialysate. Such carriers include, for example, synthetic or naturally-occurring organic macro-molecular compounds; inorganic 30 materials, such as glass beads, silica gel, alumina, and activated charcoal; and materials coated with polysaccharide(s) or synthetic polymer(s) thereof.

A carrier consisting of macromolecular compounds is exemplified by polymethyl methacrylate polymer, polyacrylonitrile polymer, 35 polysulfone polymer, vinyl polymer, polyolefin polymer, fluorine polymer, polyester polymer, polyamide polymer, polyimide polymer,

polyurethane polymer, polyacryl polymer, polystyrene polymer, polyketone polymer, silicon polymer, cellulose polymer, and chitosan polymer. More specifically, carriers are exemplified by polysaccharides, such as agarose, cellulose, chitin, chitosan, 5 sepharose, dextran, etc., and derivatives thereof; and polyester, polyvinyl chloride, polystyrene, polysulfone, polyethersulfone, polypropylene, polyvinyl alcohol, polyallylether sulfone, polyacrylic ester, polymethacrylic ester, polycarbonate, acetylated cellulose, polyacrylonitrile, polyethylene terephthalate, polyamide, 10 silicone resin, fluororesin, polyurethane, polyetherurethane, and polyacrylamide, and derivatives thereof. The macromolecular materials can be used alone or in combination of two or more kinds of macromolecules. In the latter case, a biguanide agent is immobilized on at least one of the macromolecules. Further, a 15 biguanide agent may be immobilized alone on a carrier, or two of more kinds of biguanide agents may be immobilized on the carrier.

There is no restriction on the shape of the carrier immobilizing the biguanide agent of the present invention. For example, the carrier can be membranous, fibrous, granular-shaped, hollow 20 fiber-like, non-woven fabric-like, porous, honeycomb-shaped, and so on. The area of the carrier, to be contacted with blood or dialysate, can be controlled by changing the thickness, surface area, diameter, length, shape, and/or size of the carrier.

A biguanide agent can be immobilized on the above-mentioned 25 carrier using conventional methods, such as physical adsorption, a specific biochemical binding reaction, ion binding, covalent bonding, grafting, and so on. According to needs, spacers can be inserted between carrier and biguanide agent. Preferably, a carrier and a biguanide agent are bound by covalent bonds so as to minimize the 30 amount of the biguanide agent released. Functional groups on the carrier are utilized for covalently binding a biguanide agent thereto. The functional groups include, for example, hydroxyl group, amino group, aldehyde group, carboxyl group, thiol group, silanol group, amide group, epoxy group, succinylimido group, and so on; however, 35 the functional groups of the present invention are not limited to these groups. The covalent bonds are exemplified by ester linkage,

ether linkage, amino linkage, amid linkage, sulfide linkage, imino linkage, disulfide linkage, and so on.

The carrier with a biguanide agent of the present invention immobilized on it can be sterilized by conventional sterilization methods. Specifically, sterilization methods includes, for example, autoclaving, gamma-ray irradiation, gas sterilization, and so on.

Biguanide agent-immobilized carriers can be contacted with blood in various ways. Examples thereof include: a method of trapping carbonyl compounds in patient blood by infusing the blood collected from the patient into a blood bag filled with biguanide agent-immobilized carriers; a method where blood is circulated in a cartridge column filled with bead carriers or fiber carriers, or the like, on which a biguanide agent has been immobilized; and so on. Instead of whole blood, blood plasma separated therefrom may be also treated according to the method. The treated blood may be returned to the patient or, if required, may be stored in a blood bag, or the like. Carbonyl compounds that generate and/or accumulate during storage can be also trapped by adding carriers immobilized with biguanide agents into the blood bags.

20 The contact between blood and carriers, on which a biguanide agent of this invention has been immobilized, can be carried out during the blood purification step, including hemodialysis, blood filtration, blood filtration dialysis, blood adsorption, and blood plasma separation.

25 For example, both hemodialysis and trapping of carbonyl compounds can be carried out simultaneously in hemodialysis patients by placing carriers on which a biguanide agent has been immobilized in the hemodialysis circuit. Herein, a biguanide agent is preferably immobilized on the hemodialysis membrane. Known types of dialysis membranes can be used as carriers. Examples include: cellulose derivatives, such as regenerated cellulose, and cellulose triacetate; polymethyl methacrylate; polyolefin; polysulfone; polyacrylonitrile (PAN); polyamide; polyimide; polyether nylon; silicon; polyester copolymers; and so on; however, the present invention is not limited thereto. Instead of using a dialysis membrane as a carrier, a column filled with carriers, on which a biguanide agent has been immobilized,

may be placed in the hemodialysis circuit as described above. By contacting patient's blood with carriers, on which a biguanide agent has been immobilized, carbonyl compounds are trapped from the blood, and the damaging activity of the compounds against the living body are eliminated and rendered nontoxic. Anticoagulants may be combined to prevent blood-clotting during extracorporeal circulation. Such anticoagulants include, for example, heparin, low-molecular-weight heparin, Futhan (Nafamostat mesilate), and so on. They may also be immobilized on carriers.

10 A method wherein peritoneal dialysate, instead of blood, is contacted with an immobilized biguanide agent is also useful to improve the carbonyl stress state. For example, carbonyl compounds generated and/or accumulated during storage can be trapped by enclosing the peritoneal dialysate in a container wherein a biguanide agent is immobilized therein; or in a container comprising a biguanide agent immobilized on particulates or fibrous carriers. In the latter system, the insoluble carriers can be separated from the peritoneal dialysate by filtration and the like. Alternatively, a carbonyl compound-trapping cartridge is prepared by filling a column with carrier beads or fibrous carriers, on which a biguanide agent is immobilized. Then, peritoneal dialysate is contacted with the carrier in the cartridge, and then, the fluid is infused into peritoneal cavity. When the carbonyl compound-trapping cartridge is contacted with peritoneal dialysate at the time of peritoneal infusion,

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20 although it is impossible to remove patient-derived carbonyl compounds that accumulate in the fluid during the dialysis, carbonyl compounds originally present in the dialysate can be eliminated.

25 Alternatively, when peritoneal dialysis treatment is conducted by using a closed circuit, wherein peritoneal dialysate is circulated by a small circulating pump, it is possible to remove not only carbonyl compounds originally present in the dialysate but also those that

30 accumulate in the peritoneal cavity during dialysis by installing the above-mentioned carbonyl compound-trapping cartridge containing carriers with immobilized biguanide agent within the circuit.

35 It is predicted that there may be some cases where carbonyl compounds in patient blood are not completely eliminated during

dialysis if the quantity of biguanide agent used for the contact with blood or dialysate is too small. Pre-determination of the quantity of carbonyl compounds in patient blood is particularly difficult. Thus, to be most effective, it is preferable to maintain as many 5 biguanide agents as active as possible, within a range that ensures the safety of the patient. The dose of a biguanide agent can be adjusted by altering the quantity of biguanide agent immobilized on the carriers, or the dose of carriers on which the biguanide agent has been immobilized.

10 The effect of a carbonyl stress-decreasing agent of the present invention can be confirmed by monitoring the concentration of carbonyl compounds or AGE in blood. The *in vivo* effect can be assessed by comparing blood AGE levels between a control group and a group wherein 15 a carbonyl stress-decreasing agent of the present invention has been administered. The control group may be an untreated group or a group wherein physiological saline or a control agent, consisting of the decreasing agent without the base agent, i.e. biguanide agent, has been administered. Glyoxal (GO), methylglyoxal (MGO), and 3-deoxyglucosone (3DG) can be used as indices of carbonyl compounds. 20 The levels of these carbonyl compounds can be readily determined by HPLC or the like as shown in the Examples (Ohmori S. et al., J. Chromatogr. 414:149-155, 1987; Yamada H., J. Biol. Chem. 269:20275-20280, 1994). Alternatively, the levels of the carbonyl compounds can be determined by reacting them with 25 2,4-dinitrophenylhydrazine (2,4-DNPH) under an acidic condition, and measuring the optical density of colored products of the reaction at 360nm. Further, pentosidine or the like can be used as an index of AGE. A method for quantifying pentosidine with reverse-phase HPLC is already known in the art (Miyata T, et al., J Am Soc Nephrol 7: 30 1198-1206, 1996).

Generally, orally administered biguanide agents exhibit the maximal blood level 1 to 2 hours after administration (N. Engl. J. Med. 334:574-579, 1996). Accordingly, the effect of a decreasing agent of the present invention can be tested by monitoring changes 35 of pentosidine level during this period. The effect of a decreasing agent of the invention can be assessed by decreased pentosidine levels

compared with the level of a control. On the other hand, the concentration of carbonyl compounds and AGE in blood or dialysate is detected to assess the effect of a carbonyl stress-decreasing agent of the present invention outside the living body.

5

Brief Description of the Drawings

Figure 1 depicts graphs demonstrating the carbonyl compound-trapping effect of a biguanide agent (metformin). The ordinate indicates the concentration of the carbonyl compound (μM) determined by high-performance liquid chromatography; the abscissa indicates the incubation time (hr.).

Figure 2 depicts graphs demonstrating the carbonyl compound-trapping effect of a biguanide agent (buformin). The ordinate indicates the concentration of the carbonyl compound (μM) determined by high-performance liquid chromatography; the abscissa indicates the incubation time (hr.).

Figure 3 depicts graphs demonstrating the carbonyl compound-trapping effect of a biguanide agent (phenformin). The ordinate indicates the concentration of the carbonyl compound (μM) determined by high-performance liquid chromatography; the abscissa indicates the incubation time (hr.).

Figure 4 depicts graphs demonstrating the changes of concentration of carbonyl compounds, which was determined in the absence of any biguanide agent (control). The ordinate indicates the concentration of the carbonyl compound (μM) determined by high-performance liquid chromatography; the abscissa indicates the incubation time (hr.).

Figure 5 depicts a graph demonstrating the dicarbonyl compound (glyoxal)-trapping action of biguanide agents in peritoneal dialysate. The ordinate indicates the residual rate (%) of glyoxal in the peritoneal dialysate; the abscissa indicates the type of the biguanide agent.

Figure 6 depicts a graph demonstrating the dicarbonyl compound (methylglyoxal)-trapping action of biguanide agents in peritoneal dialysate. The ordinate indicates the residual rate (%) of methylglyoxal in the peritoneal dialysate; the abscissa indicates

the type of the biguanide agent.

Figure 7 depicts a graph demonstrating the dicarbonyl compound (3-deoxyglucosone)-trapping action of biguanide agents in peritoneal dialysate. The ordinate indicates the residual rate (%) of 3-deoxyglucosone in the peritoneal dialysate; the abscissa indicates the type of the biguanide agent.

Figure 8 depicts graphs demonstrating the dicarbonyl compound (glyoxal and methylglyoxal)-trapping action of biguanide agents in peritoneal dialysis effluent. The ordinate indicates the residual rates (%) of glyoxal and methylglyoxal in the peritoneal dialysis effluent; the abscissa indicates the type and the concentration of the biguanide agent.

Figure 9 depicts graphs demonstrating the dicarbonyl compound (glyoxal and methylglyoxal)-trapping action of biguanide agents in serum. The ordinate indicates the residual rates (%) of glyoxal and methylglyoxal in the serum; the abscissa indicates the type and the concentration of the biguanide agent.

Best Mode for Carrying out the Invention

The present invention is illustrated in detail below with reference to Examples.

[Example 1] Carbonyl compound-trapping effect of biguanide agents

100 μ l of a mixed solution of representative carbonyl compounds, i.e. glyoxal (GO), methylglyoxal (MGO), and 3-deoxyglucosone (3DG) (1 mM each), was combined with 800 μ l of 0.1M phosphate buffer (pH7.4) and 100 μ l of 30mM biguanide agent solution, and was incubated at 37°C. Metformin, buformin, and phenformin were used as biguanide agents. After the incubation, the concentrations of glyoxal, methylglyoxal, and 3-deoxyglucosone in the solution were determined by high-performance liquid chromatography. More specifically, 40 μ l of 2M perchloric acid, 40 μ l of 1% o-phenylenediamine, 100 μ l of 50 μ M 2,3-butanedione were added to 100 μ l of the sample after the incubation; the mixture was stirred and incubated at 25°C for 1 hour. Quinoxaline derivatives, which are products of the reaction between dicarbonyl compounds and o-phenylenediamine, were quantified by HPLC

using a reverse-phase column according to the method of Ohmori et al. (Ohmori S. et al., J. Chromatogr. 414:149-155, 1987). Glyoxal, methylglyoxal, and 3-deoxyglucosone were used as standard samples.

The results are shown in Figure 1 (metformin), Figure 2 (buformin), and Figure 3 (phenformin). Almost no change in the carbonyl compound levels during the incubation is observed with the carbonyl compounds alone (Figure 4; control). On the other hand, in the presence of a biguanide agent selected from those mentioned above, the concentrations of glyoxal and methylglyoxal were markedly decreased depending on the incubation period by each agent, and the carbonyl compound-trapping effect of the biguanide agents was confirmed. Thus, the carbonyl compounds were verified to lose their reactivity by reacting with the biguanide agents.

15 [Example 2] Dicarbonyl-trapping action of biguanide agents in peritoneal dialysates

Peritoneal dialysates comprise carbonyl compounds generated during autoclaving or storage. The carbonyl compounds, which are generated during the processes of production or storage of the 20 peritoneal dialysate, can be preventively detoxified by previously adding a biguanide agent to the peritoneal dialysate. Alternatively, the carbonyl compounds can be removed by contacting the peritoneal dialysate with a carrier immobilized with the biguanide agent.

First, the quantities of various carbonyl compounds in a 25 commercially available peritoneal dialysate (Baxter Ltd.; Dianeal PD-4 1.5TM) were determined according to the method of Ohmori et al. (Ohmori S. et al., J. Chromatogr., 414:149-155, 1987). The concentrations of glyoxal, methylglyoxal, and 3-deoxyglucosone in the peritoneal dialysate were determined to be 11 μM, 3.5 μM, and 42 30 μM, respectively.

Then, 100 μL of a biguanide solution (dissolved in 0.1M phosphate buffer (pH 7.4)) was added to 900 μL of the peritoneal dialysate (Baxter Ltd.; Dianeal PD-4 1.5TM) to prepared a sample solution. The final concentrations of the biguanide agents were 35 adjusted to 0, 1, 5, and 10mM. Metformin, buformin, and phenformin were used as biguanide agents. The mixed sample solution was

incubated at 37°C for 4 hours.

The quantities of residual glyoxal, methylglyoxal, and 3-deoxyglucosone in the sample solution were determined by high-performance liquid chromatography (HPLC) following the 5 incubation. More specifically, 40 µl of 2M perchloric acid, 40 µl of 1% o-phenylenediamine, and 100 µl of 25 µM 2,3-butanedione were added to 100 µl of the sample after the incubation; the mixture was stirred and incubated at 25°C for 1 hour. Quinoxaline derivatives produced in the reaction of the dicarbonyl compounds and 10 o-phenylenediamine were quantified by HPLC using a reverse-phase column according to the method of Ohmori et al. (Ohmori S. et al., J. Chromatogr. 414: 149-155, 1987). Glyoxal, methylglyoxal, and 3-deoxyglucosone were used as standard samples.

The results are shown in Figure 5 (glyoxal), Figure 6 15 (methylglyoxal), and Figure 7 (3-deoxyglucosone). The quantity of each carbonyl compound is represented by the residual rate (%), taking the quantity in the absence of any biguanide agent as 100%. In the presence of a biguanide agent, the concentrations of glyoxal and methylglyoxal were decreased depending on the biguanide concentration, 20 and thus, the dicarbonyl compound-trapping effect in a peritoneal dialysate was confirmed.

[Example 3] Dicarbonyl-trapping action of a biguanide agent in peritoneal dialysis effluent

25 During peritoneal dialysis, highly reactive carbonyl compounds, along with circulating waste products, are transferred across the peritoneum and accumulated in peritoneal dialysate in the patient's peritoneal cavity. Therefore, in order to study the ability of a biguanide agent to trap carbonyl compounds in peritoneal dialysate 30 in peritoneal cavity during peritoneal dialysis, experiments were carried out to determine the effect of a biguanide agent to trap dicarbonyl compounds in peritoneal dialysis discharged from peritoneal cavity of a peritoneal dialysis patient.

First, 100 µl of biguanide solution (dissolved in 0.1M phosphate 35 buffer (pH 7.4)) was added to 900 µl of peritoneal dialysis effluent (liquid allowed to dwell in the peritoneal cavity for 2 hours) from

a peritoneal dialysis patient administered with a commercially available peritoneal dialysate (Baxter Ltd.; Dianeal PD-4 1.5TM); the final concentrations of the biguanides were 0, 5, 25, and 50mM. The mixture was incubated at 37°C for 4 hours. Metformin, buformin, and phenformin were used as biguanide agents. After the incubation, 200 µl of water and 100 µl of 2M perchloric acid were added to 200 µl of the sample. The mixture was centrifuged and filtered with a 0.45-µm filter. 50 µl of 10 µM 2,3-butanedione and 20 µl of 1% O-phenylenediamine were added to 150 µl of the filtrate. The mixture was stirred and incubated at room temperature for 2 hours. Then, 40 µl of 5N sodium hydroxide solution was added, and the mixture was stirred. 600 µl of toluene was added to the mixture for extraction. Further, 2M perchloric acid was added to 500 µl of the toluene phase, and the mixture was stirred to extract quinoxaline derivatives. To 10 quantify glyoxal and methylglyoxal, the aqueous phase was analyzed by HPLC using a reverse-phase column according to the method of Ohmori et al. (Ohmori S. et al., J. Chromatogr. 414: 149-155, 1987). Glyoxal and methylglyoxal were used as standard samples.

The results are shown in Figure 8. The quantity of each carbonyl compound is represented by the residual rate (%), taking the quantity in the absence of any biguanide agent as 100%. In the presence of a biguanide agent, the concentrations of glyoxal and methylglyoxal decreased, depending on the biguanide concentration, and thus, the dicarbonyl compound-trapping effect in a peritoneal dialysis effluent was confirmed.

[Example 4] The action of a biguanide agent to trap dicarbonyl compounds in serum

In order to study the effect of the carbonyl stress-decreasing agents of the present invention in patients treated with blood dialysis, the action of biguanide agents to trap dicarbonyl compounds in serum was examined.

First, 100 μ l of biguanide solution (dissolved in 0.1M phosphate buffer (pH 7.4)) was added to 900 μ l serum from a blood dialysis patient; the final concentrations of the biguanides were 0, 5, 25, and 50mM. The mixture was incubated at 37°C for 4 hours. Metformin,

buformin, and phenformin were used as biguanide agents. Glyoxal and methylglyoxal were quantified by the same method as in Example 3.

The results are shown in Figure 9. The quantity of each carbonyl compound is represented by the residual rate (%), taking the quantity 5 in the absence of any biguanide agent as 100%. In the presence of a biguanide agent, the concentrations of glyoxal and methylglyoxal decreased, and thus, the dicarbonyl compound-trapping effect in serum was confirmed.

10 Industrial Applicability

The present invention provides carbonyl stress-decreasing agents that are expected to remove carbonyl compounds both within a living body (*in vivo*) and outside the body (*ex vivo*). Biguanide agents constituting the carbonyl stress-decreasing agents of the 15 present invention are oral antidiabetic drugs that have been used clinically, and therefore can be administered orally or by other routes for administration. The biguanide agents can be used as agents having direct action against *in vivo* carbonyl stress. Furthermore, the agents can be provided conveniently due to highly sophisticated 20 formulation techniques previously established for the agents.

CLAIMS

1. A carbonyl stress-decreasing agent comprising a biguanide agent or pharmacologically acceptable salt thereof as an active ingredient.
 2. The carbonyl stress-decreasing agent of claim 1, wherein the biguanide agent is a compound selected from the group consisting of: phenformin, metformin, buformin, and pharmacologically acceptable salts thereof.
 3. A carrier, on which a biguanide agent has been immobilized.
 4. An adsorbent of carbonyl compounds comprising the carrier of claim 3.
 5. A method for removing carbonyl compounds comprising the step of contacting the carrier of claim 3 with a body fluid selected from the group consisting of: blood, blood plasma and peritoneal dialysate.

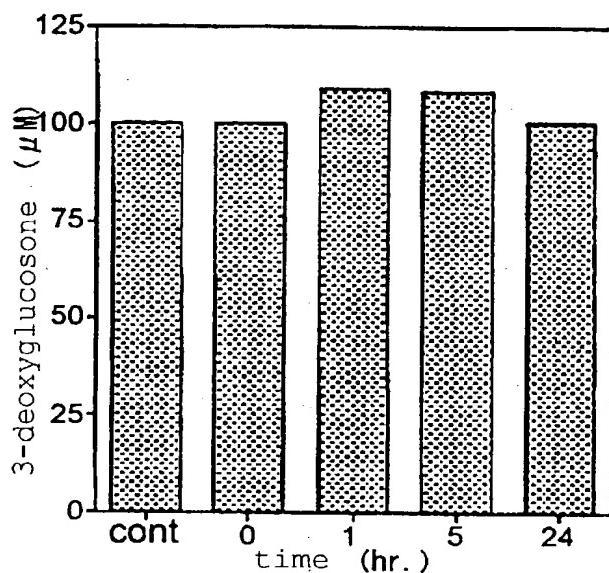
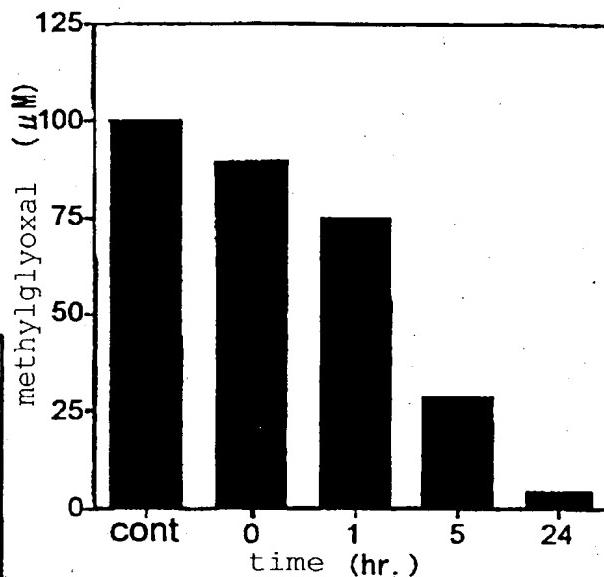
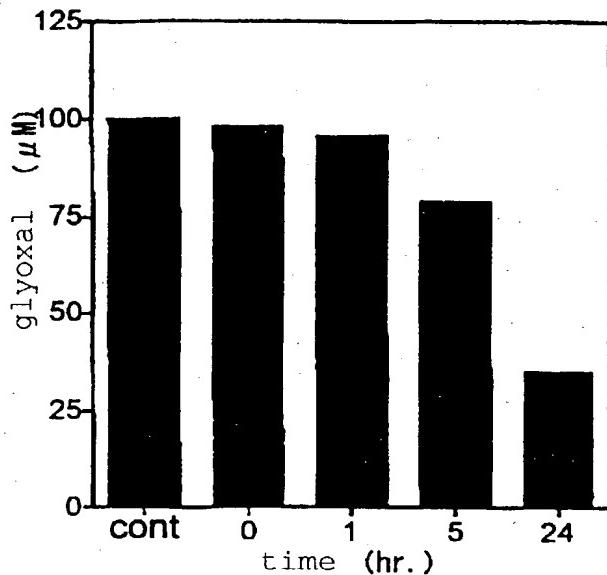
ABSTRACT

The present invention provides agents that decrease carbonyl stress comprising biguanides, such as metformin. Through oral 5 administration and such, the carbonyl stress-decreasing agents of the present invention are useful as drugs directly acting against carbonyl stress *in vivo*.

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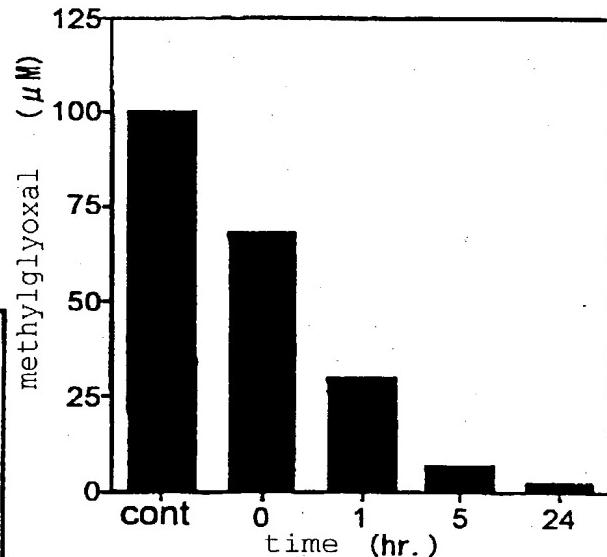
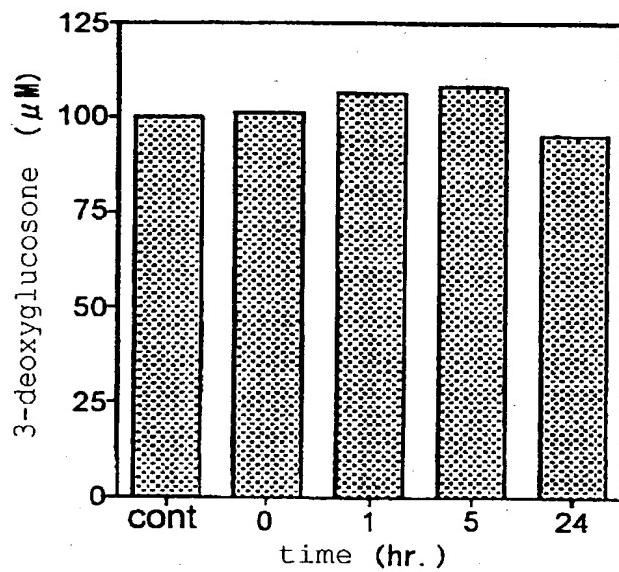
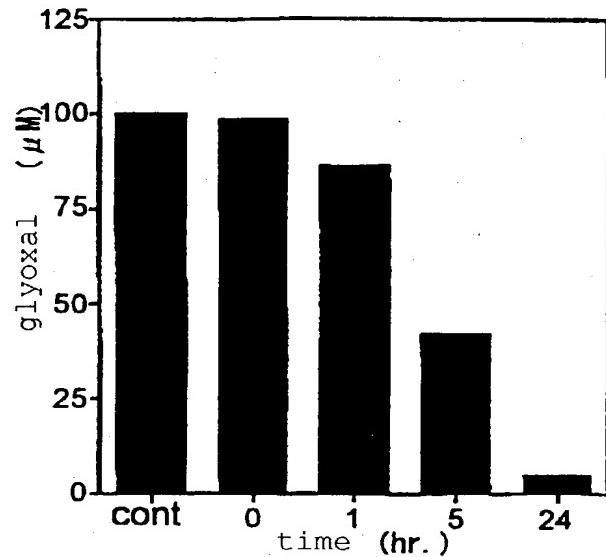
Figure 1



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Figure 2

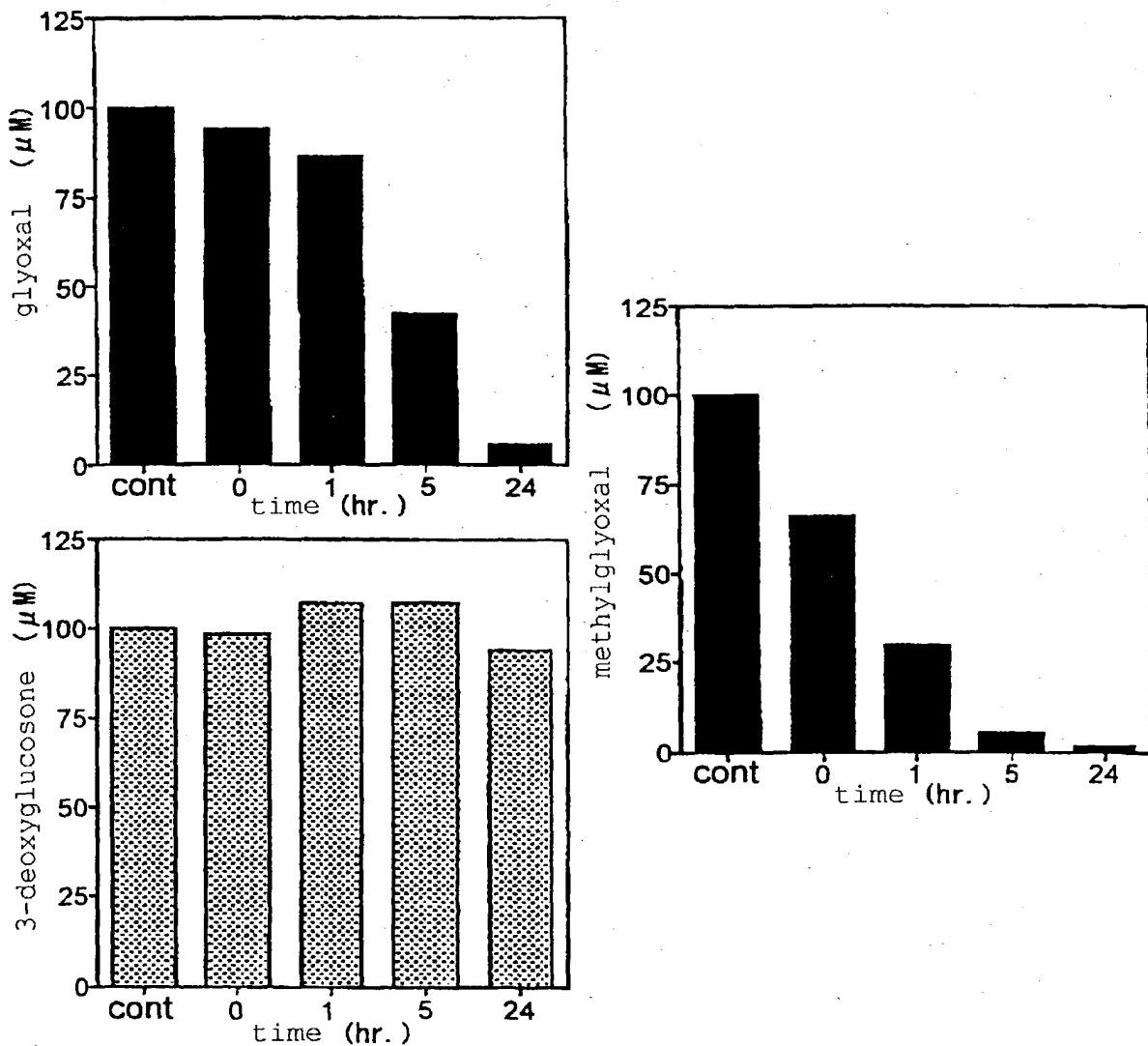


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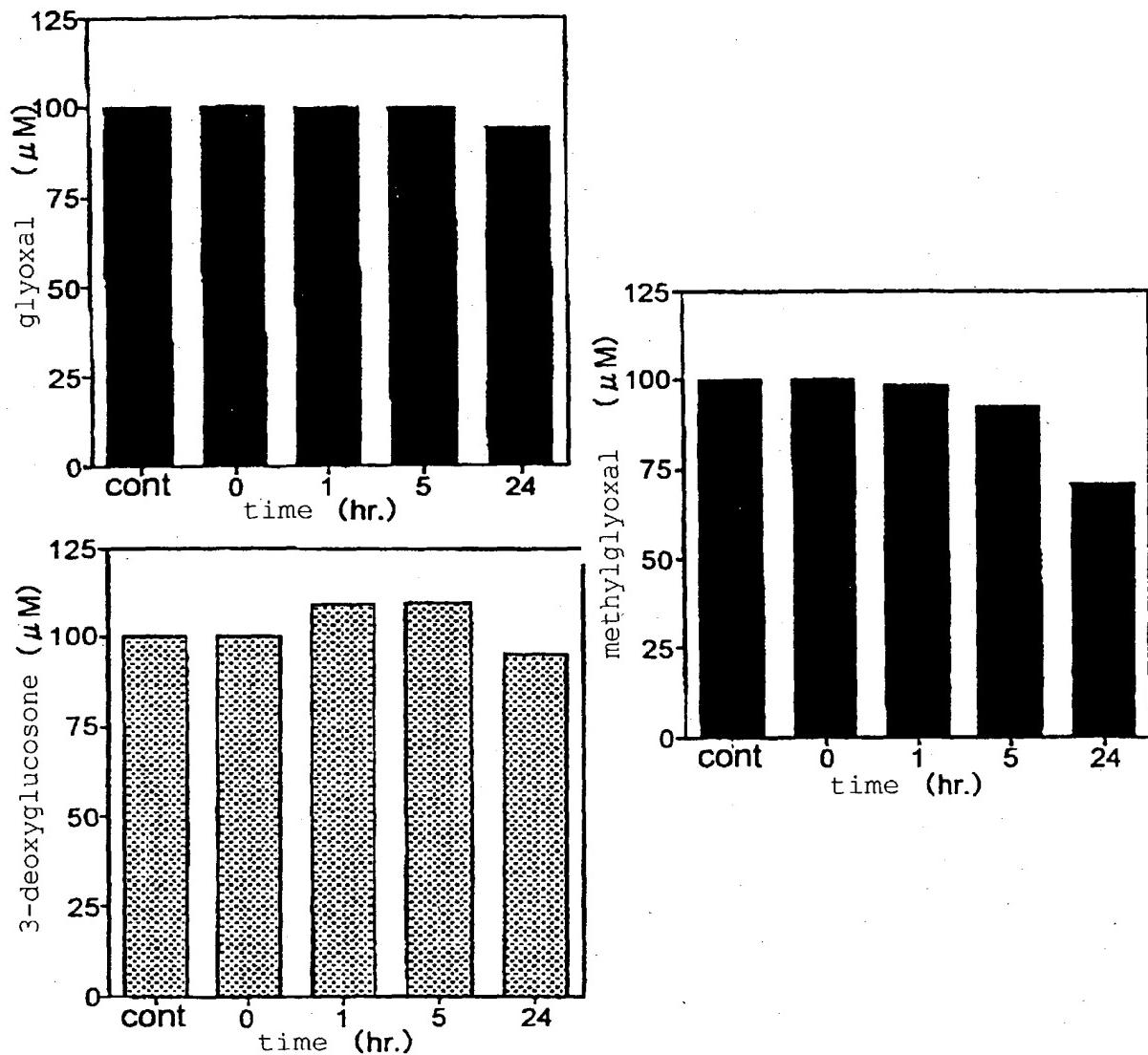
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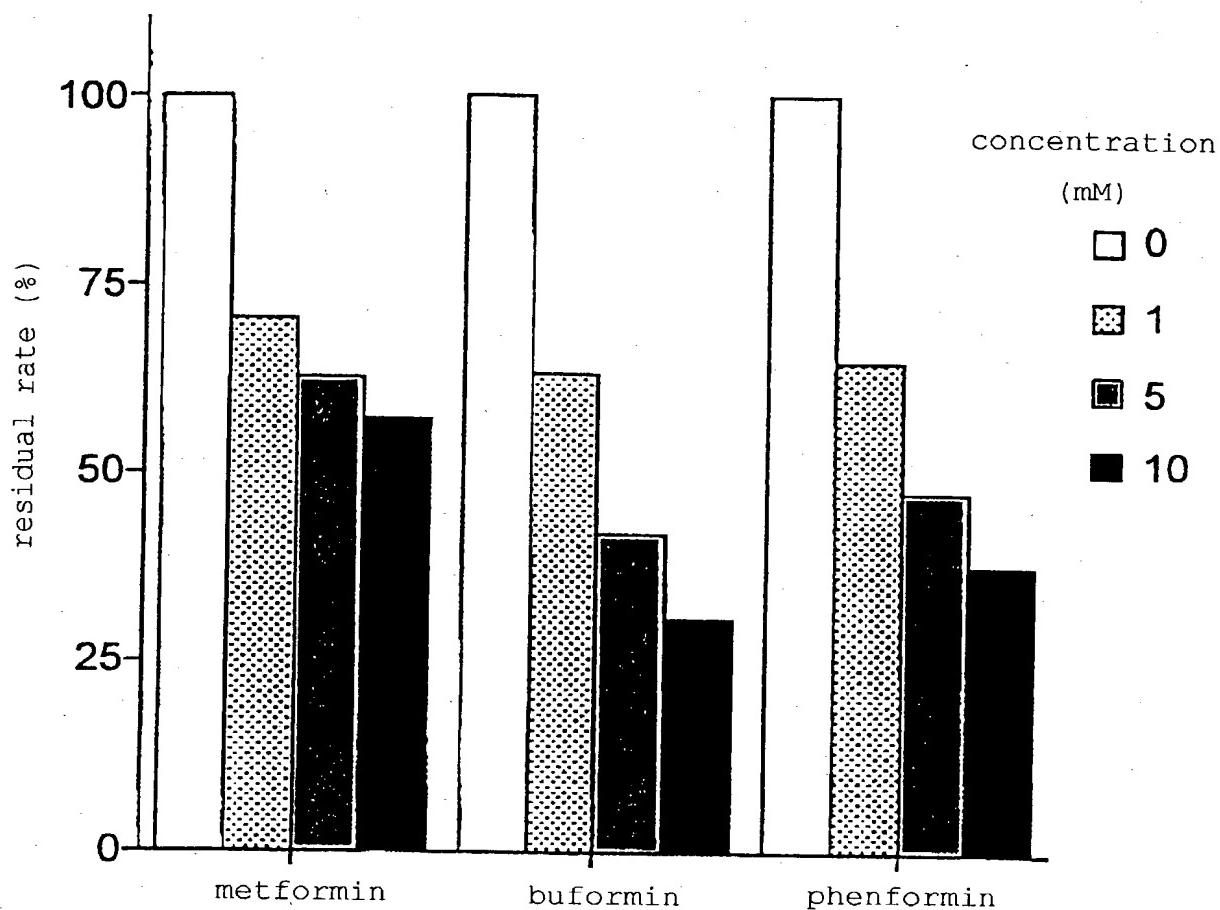
Figure 4



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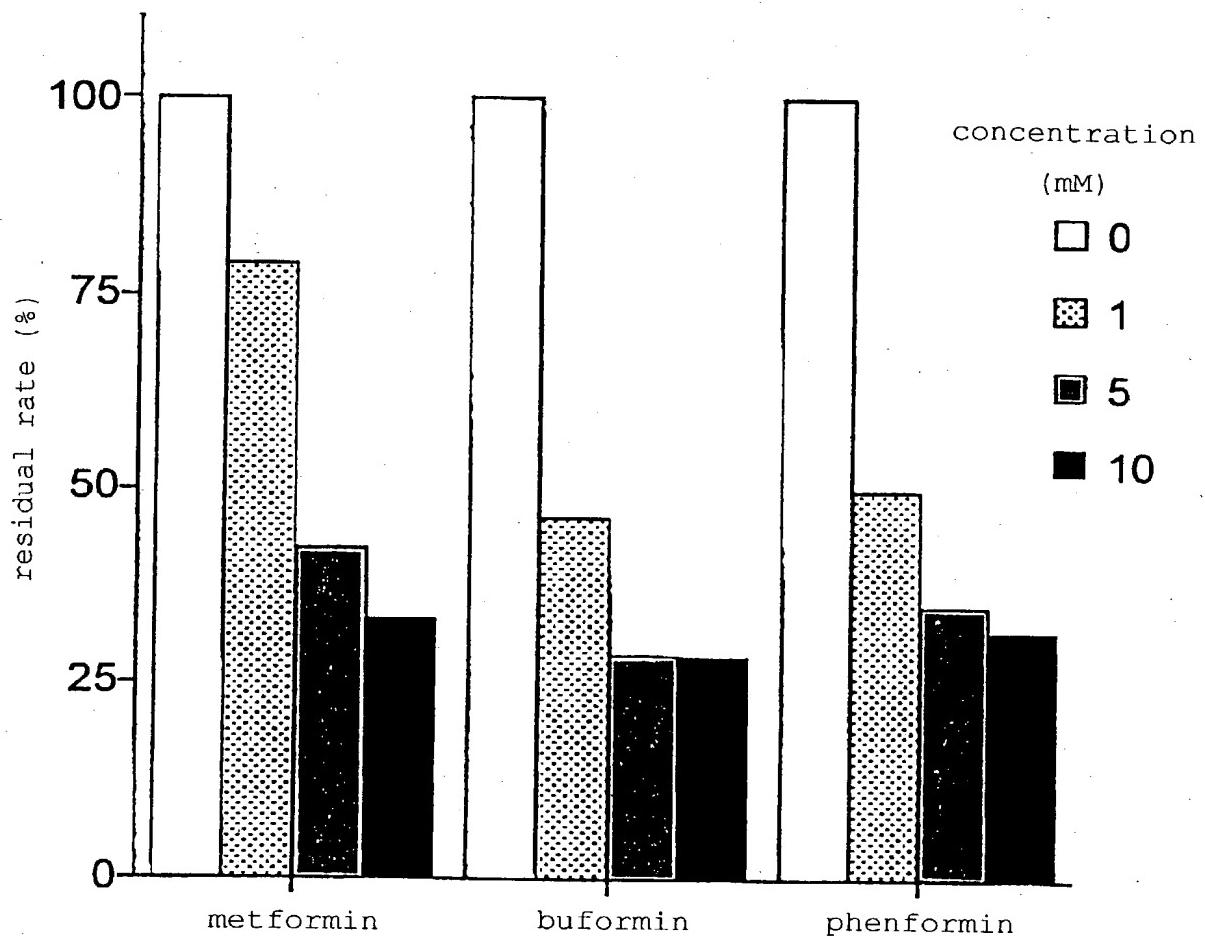
Figure 5



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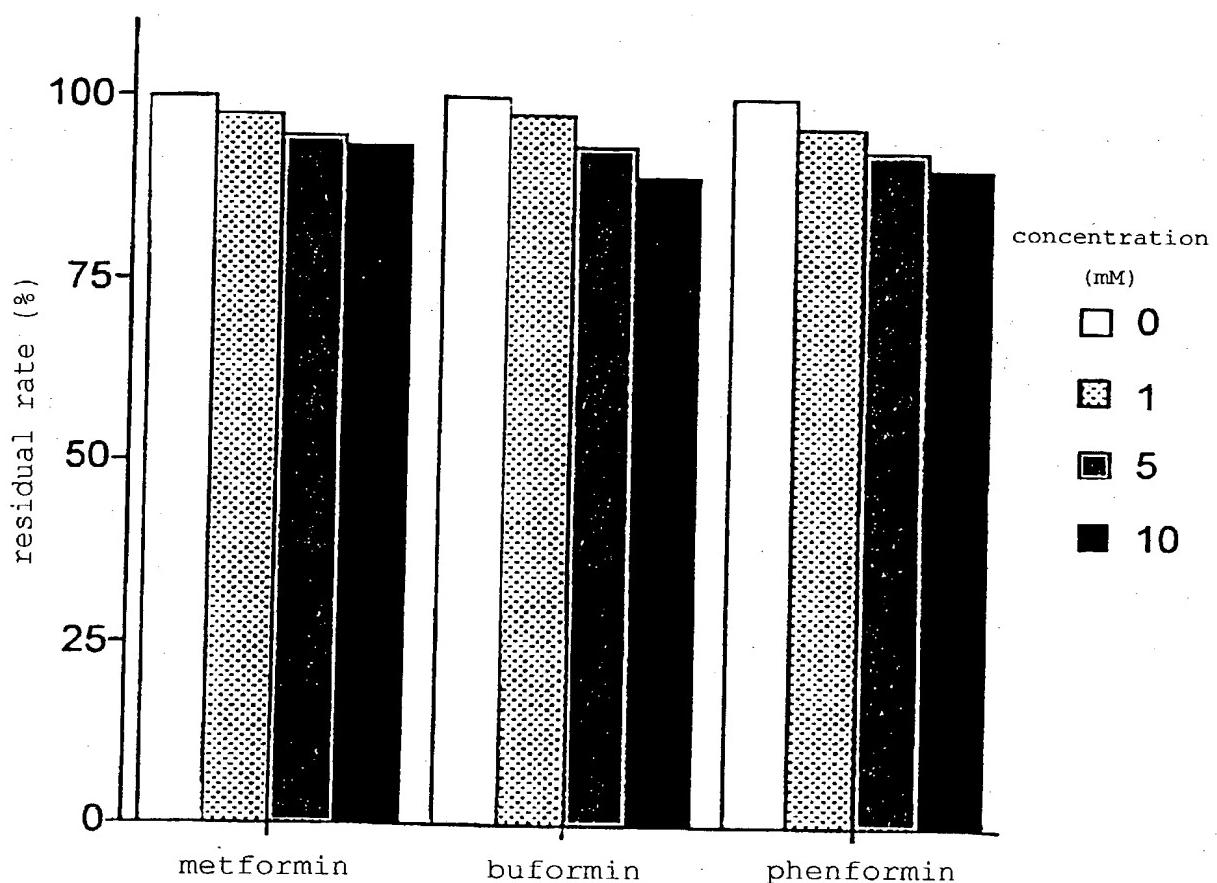
Figure 6



INVESTIGATIONS - DRUGS
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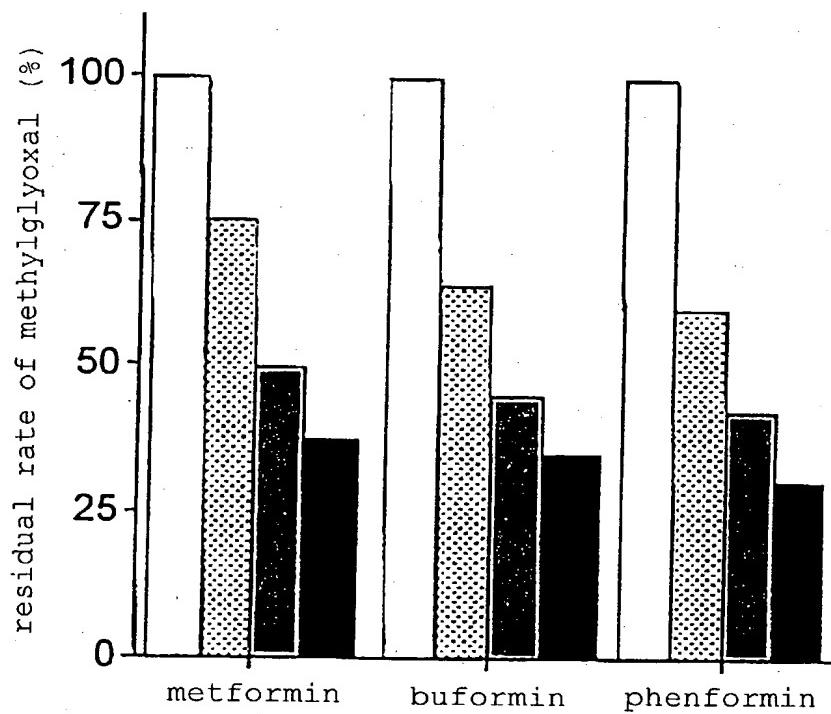
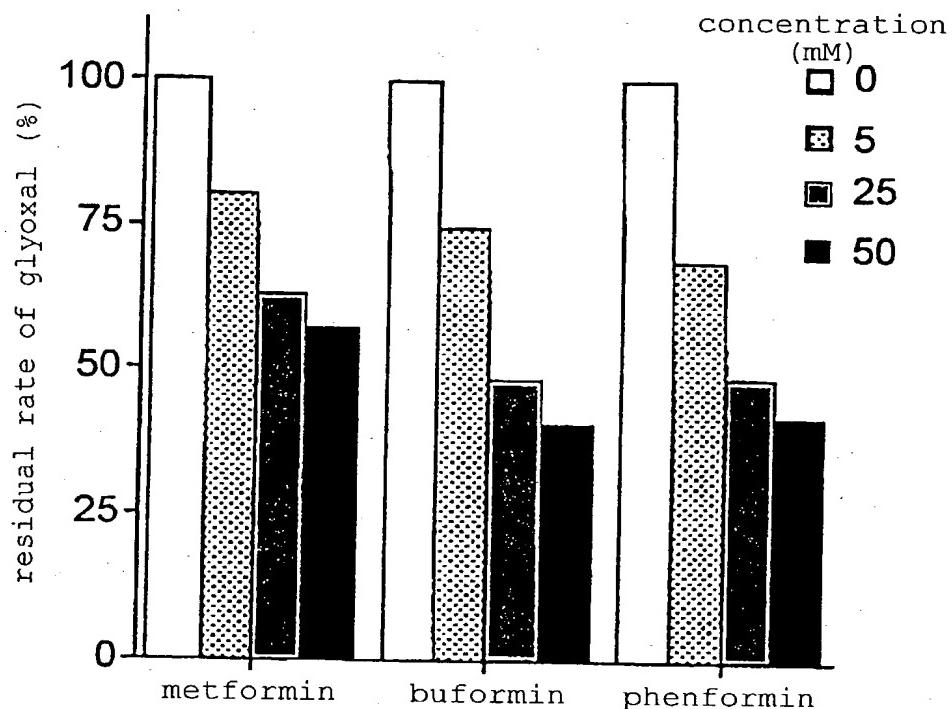
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Figure 7



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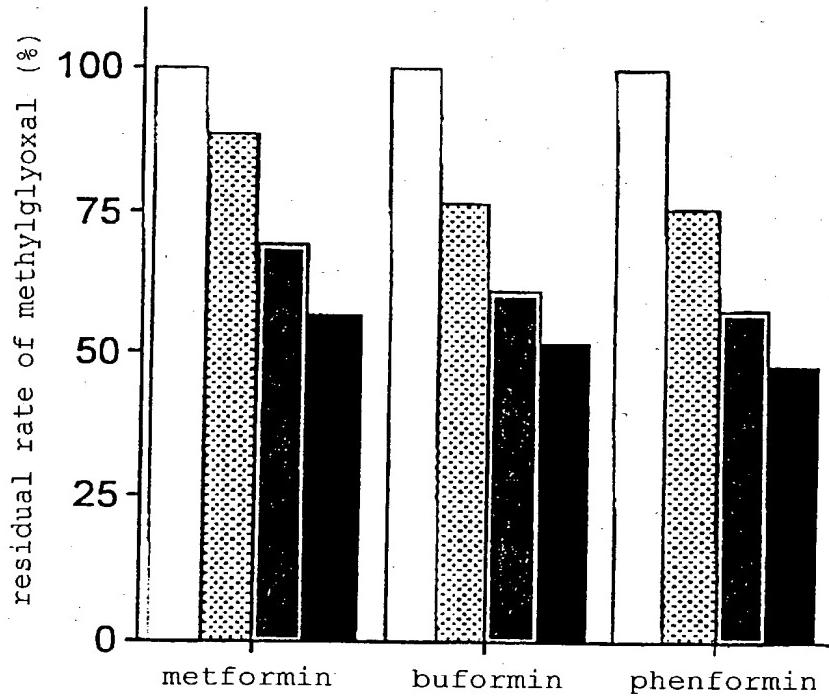
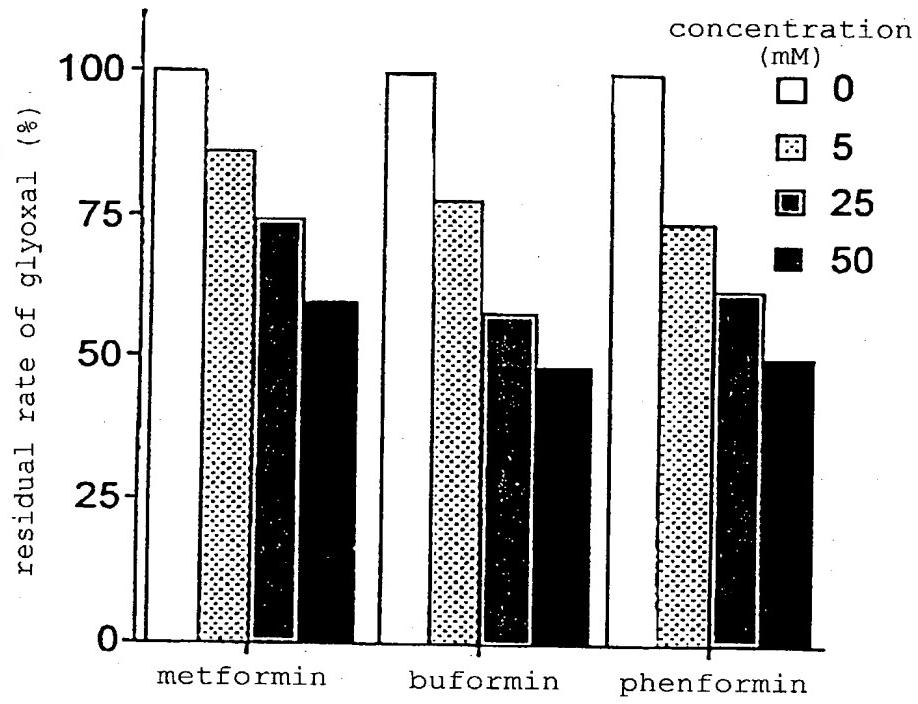
Figure 8



7/089789

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Figure 9



COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY (Includes Reference to PCT International Applications)

ATTORNEY'S DOCKET NO.
SHIM1130

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

CARBONYL STRESS-DECREASING AGENT

the specification of which (check only one item below):

is attached hereto.

was filed as United States application
Serial No. 10/089,789

on

International Filing Date: 6 October 2000
and was amended
on _____ (if applicable).

was filed as PCT international application
Number PCT/JP00/06987

on October 6, 2000

and was amended under PCT Article 19

on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations §1.56(a).

I hereby claim foreign priority benefit under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S) AND ANY PRIORITY CLAIMS UNDER 35 U.S.C. 119:

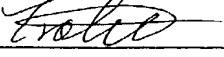
COUNTRY (If PCT indicate "PCT")	APPLICATION NUMBER	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC 119
Japan	11/285735	6 October 1999	<input checked="" type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO



COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY (CONTINUED) (Includes Reference to PCT International Applications)				ATTORNEY'S DOCKET NO. SHIM1130
<p>I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application(s) and the national or PCT international filing date of this application.</p>				
PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 35 U.S.C. 120:				
U.S. APPLICATIONS		STATUS (MARK ONE)		
U.S. APPLICATION NUMBER		U.S. FILING DATE		PATENTED PENDING ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.				
PCT APPLICATION NO.	PCT FILING DATE	U.S. SERIAL NUMBERS ASSIGNED (if any)		
PCT/JP00/06987	6 October 2000		X	
POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith. (List name and registration no.)				
CUSTOMER NUMBER	28,213	PLACE CUSTOMER LABEL HERE		
SEND CORRESPONDENCE TO: Lisa A. Haile, Reg. No. 38,347 GRAY CARY WARE & FREIDENRICH 4365 Executive Drive, Suite 1100 San Diego, CA 92121-2133			DIRECT TELEPHONE CALLS TO: <u>LISA A. HAILE</u> Telephone: (858) 677-1456 Fax: (858) 677-1465	
2 0 1	FULL NAME OF INVENTOR <u>Toshio Miyata</u>	FAMILY NAME	FIRST GIVEN NAME Toshio	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY Kanagawa	STATE OR FOREIGN COUNTRY Japan <u>JPX</u>	COUNTRY OF CITIZENSHIP Japan
	POST OFFICE ADDRESS	POST OFFICE ADDRESS 102 Ekuseru Isenhara, 16-25, Sakuradai 2-chome, Isenhara-shi	CITY Kanagawa	STATE & ZIP CODE/COUNTRY 259-1132 JAPAN
2 0 2	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY
2 0 3	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY
2 0 4	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY

ADDITIONAL INVENTOR INFORMATION ATTACHED

I hereby declare that all statements made herein are of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any issuing thereon.

SIGNATURE OF INVENTOR 201	SIGNATURE OF INVENTOR 202	SIGNATURE OF INVENTOR 203
		
DATE: June 18, 2002	DATE:	DATE::